Amendments to the Claims:

Please amend claims 1, 3, 6, 7, 10-13, 34-36 and 38-41, cancel claims 2, 4-5, and 14-33, and add new claims 42-48. This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

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- 1 (currently amended): A method for modulating reducing expression of a mammalian SREBP-1 gene by administering a modulator compound that promotes or an antagonist of LXR α that inhibits LXR α -mediated expression of the SREBP-1 gene to a cell that comprises an SREBP-1 gene and an LXR α polypeptide, wherein said antagonist is an oxysterol.
- 1 2 (canceled).
- 3 (currently amended): The method of claim 1, wherein the modulator
 compound promotes or antagonist inhibits LXRα-mediated expression of the SREBP-1c
 transcript.
- 1 4 (canceled).
- 1 5 (canceled).
- 6 (currently amended): The method of claim § 1, wherein the cell further
 comprises one or more genes that encode an enzyme involved in fatty acid and triglyceride
 metabolism and contacting the cell with the modulator compound antagonist inhibits expression
 of one or more of the genes that encode an enzyme involved in fatty acid and triglyceride
 metabolism.
- 7 (currently amended): The method of claim 1 6, wherein the enzyme involved in fatty acid and triglyceride metabolism is selected from the group consisting of fatty acid synthase, acetyl CoA carboxylase, steroyl CoA desaturase, and lipoprotein lipase.

Appl. No. 09/848,990 Amdt. dated January 24, 2005 Reply to Office Action of September 24, 2004

1	8 (original): The method of claim 1, wherein the cell is in a mammal.
1	9 (original): The method of claim 8, wherein the mammal is a human.
1	10 (currently amended): The method of claim 8, wherein the modulator
2	compound is an antagonist of LXRα and triglyceride levels in the mammal are reduced.
1	11 (currently amended): The method of claim 8, wherein the modulator
2	compound is an antagonist of LXR and insulin levels in the mammal are reduced.
1	12 (currently amended): A method of modulating triglyceride levels in a
2	mammal, the method comprising administering to the mammal an effective amount of a
3	modulator compound that comprises at least one of LXRα antagonist and LXRα agonist activity
4	that promotes or inhibits LXR\alpha-mediated expression of an SREBP-1 gene in cells of the
5	mammal, wherein the modulator compound is an oxysterol.
1	13 (currently amended): The method of claim 12, wherein the mammal is a
2	human.
1	14-33 (canceled)
1	34 (currently amended): A method for ameliorating a condition associated with
2	abnormal abnormally high SREBP-1 expression in a mammal, the method comprising
3	administering to the mammal a therapeutically effective amount of a LXR\alpha antagonist, wherein
4	said antagonist is an oxysterol.
1	35 (currently amended): The method of claim 34, wherein the condition
2	associated with abnormal abnormally high SREBP-1 expression is hypertriglyceridemia.
1	36 (currently amended): The method of claim 34, wherein the condition
2	associated with abnormal abnormally high SREBP-1 expression is lipodystrophy.

Appl. No. 09/848,990 Amdt. dated January 24, 2005 Reply to Office Action of September 24, 2004

1	37 (original): The method of claim 36, wherein the lipodystrophy is congenital
2	generalized lipodystrophy.
1	38 (currently amended): The method of claim 34, wherein the condition
2	associated with abnormal abnormally high SREBP-1 expression is insulin resistance.
1	39 (currently amended): The method of claim 34, wherein the condition
2	associated with abnormal abnormally high SREBP-1 expression is an elevated plasma insulin
3	level.
1	40 (currently amended): The method of claim 34, wherein the condition
2	associated with abnormal abnormally high SREBP-1 expression is hyperglycemia and/or
3	diabetes mellitus.
1	41 (currently amended): The method of claim 34, wherein the condition
2	associated with abnormal abnormally high SREBP-1 expression is a syndrome associated with
3	treatment of AIDS by administration of an HIV protease inhibitor, which syndrome is
4	characterized by one or more of lipodystrophy, insulin resistance and hyperlipidemia.
1	42. (new): The method of claim 34, wherein the condition associated with
2	abnormally high SREBP-1 expression is pancreatitis.
1	43. (new): The method of claim 12, wherein the modulator compound is an
2	agonist of LXR α and promotes LXR α -mediated expression of the SREBP-1 gene.
1	44 (new): The method of claim 12, wherein the modulator compound promotes
2	or inhibits LXR α -mediated expression of the SREBP-1c transcript.
1 ·	45 (new): The method of claim 43, wherein the modulator compound is 24,25-
2	epoxycholesterol.

Appl. No. 09/848,990 Amdt. dated January 24, 2005 Reply to Office Action of September 24, 2004

46 (new): The method of claim 12, wherein the modulator compound is an 1 antagonist of LXR α and inhibits LXR α -mediated expression of the SREBP-1 gene. 2 47. (new): A method of increasing triglyceride levels in a mammal, the method 1 comprising administering to the mammal an effective amount of an agonist of LXR α that 2 promotes LXR α -mediated expression of an SREBP-1 gene in cells of the mammal. 3 48. (new): The method of claim 47, wherein the agonist is selected from the 1 group consisting of an oxysterol, N-methyl-N-[4-(2,2,2-trifluoro-l-hydroxy-1-trifluoromethyl-2 ethyl)-phenyl]-benzenesulfonamide (T0314407), N-(2,2,2-trifluoro-ethyl)-N-[4-(2,2,2-trifluoro-l-3 hydroxy-1-trifluoromethyl-ethyl)-phenyl]-benzenesulfonamide (T0901317), and mixtures 4 5 thereof. 49. (new): A method of decreasing triglyceride levels in a mammal, the method 1 comprising administering to the mammal an effective amount of an antagonist of LXR\alpha that 2 inhibits LXRa-mediated expression of an SREBP-1 gene in cells of the mammal, wherein the 3 modulator compound is an oxysterol. 4